

FORM PTO-1399 (Modified) (REV 11-99)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTORNEY'S DOCKET NUMBER 00537-165002
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371			U.S. APPLICATION NO. (IF KNOWN, SEE 35 CFR 09/600648)
INTERNATIONAL APPLICATION NO. PCT/US99/01180	INTERNATIONAL FILING DATE 20 January 1999	PRIORITY DATE CLAIMED 29 January 1998	
TITLE OF INVENTION ABSORBABLE MICROPARTICLES			
APPLICANT(S) (FOR DO/EO/US) SHALABY, Shalaby Wahba			
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:			
<ol style="list-style-type: none"> <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371. <input checked="" type="checkbox"/> This is an express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1). <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371 (c) (2)) <ol style="list-style-type: none"> <input type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau). <input type="checkbox"/> has been transmitted by the International Bureau. <input checked="" type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US). <input type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)). <input type="checkbox"/> A copy of the International Search Report (PCT/ISA/210). <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3)) <ol style="list-style-type: none"> <input type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau). <input type="checkbox"/> have been transmitted by the International Bureau. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. <input checked="" type="checkbox"/> have not been made and will not be made. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)). <input type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)). <input type="checkbox"/> A copy of the International Preliminary Examination Report (PCT/IPEA/409). <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)). 			
Items 13 to 20 below concern document(s) or information included:			
<ol style="list-style-type: none"> <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included. <input type="checkbox"/> A FIRST preliminary amendment. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment. <input type="checkbox"/> A substitute specification. <input type="checkbox"/> A change of power of attorney and/or address letter. <input checked="" type="checkbox"/> Certificate of Mailing by Express Mail <input type="checkbox"/> Other items or information: 			
"Express Mail" label number: EL445347799US Date of Deposit: 20 July 2000 I hereby certify that under 37 CFR 1.10 that this correspondence is being deposited with the United States Postal Service as "Express Mail Post Office To Addressee" with sufficient postage on the date indicated above and is addressed to the Assistant Commissioner for Patents, Washington, D.C. 20231. <i>July 20, 2000</i> <i>Samantha Bell</i> <i>Samantha Bell</i>			

09/600648

PCT/US99/01180

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21. The following fees are submitted:

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :

- ☐ Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO \$970.00
- ☒ International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$840.00
- ☐ International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$690.00
- ☐ International preliminary examination fee paid to USPTO (37 CFR 1.482) but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$670.00
- ☐ International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4) \$96.00

ENTER APPROPRIATE BASIC FEE AMOUNT =

\$840.00

Surcharge of \$130.00 for furnishing the oath or declaration later than months from the earliest claimed priority date (37 CFR 1.492 (e)). ☐ 20 ☐ 30

\$0.00

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	
Total claims	46 - 20 =	26	x \$18.00	\$468.00
Independent claims	2 - 3 =	0	x \$78.00	\$0.00
Multiple Dependent Claims (check if applicable)			<input type="checkbox"/>	\$0.00

TOTAL OF ABOVE CALCULATIONS =

\$1,308.00

Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28) (check if applicable). ☐

\$0.00

SUBTOTAL =

\$1,308.00

Processing fee of \$130.00 for furnishing the English translation later than months from the earliest claimed priority date (37 CFR 1.492 (f)). ☐ 20 ☐ 30 +

\$0.00

TOTAL NATIONAL FEE =

\$1,308.00

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) (check if applicable). ☐

\$0.00

TOTAL FEES ENCLOSED =

\$1,308.00

Amount to be:

refunded \$

charged \$

- ☒ A check in the amount of **\$1,308.00** to cover the above fees is enclosed.
- ☐ Please charge my Deposit Account No. _____ in the amount of _____ to cover the above fees.
A duplicate copy of this sheet is enclosed.
- ☒ The Commissioner is hereby authorized to charge any fees which may be required, or credit any overpayment to Deposit Account No. **06-1050** A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

Y. Rocky Tsao
Fish & Richardson P.C.
225 Franklin Street
Boston, Massachusetts 02110-2804
United States of America

SIGNATURE

Y. Rocky Tsao

NAME

34,053

REGISTRATION NUMBER

DATE

534 Rec'd PCT/PTC 20 JUL 2000

ABSORBABLE MICROPARTICLESBackground of the Invention

5 This invention pertains to a sustained release complex of one or more peptide, one or more protein or a combination thereof immobilized on an absorbable polymer microparticle optionally having an absorbable polymer
10 coating. The microparticle complex of this invention comprises a peptide(s) and/or protein(s) which have at least one amino group and/or at least one carboxyl group per molecule and a solid absorbable polyester microparticle having surface and subsurface carboxylic groups or amino
15 groups in sufficient amounts to bind the peptide(s) and/or protein(s) so that the immobilized peptide(s) or protein(s) represent 0.1% to 30% of the total mass of the microparticle complex. The microparticle complex with immobilized peptide(s) and/or protein(s) are optionally
20 further encased individually or in groups with an absorbable polymer to control, further, the release of the immobilized peptide(s) and/or protein(s). To control the release of the immobilized peptide(s) and/or protein(s) even further, the encased microparticles can be
25 incorporated into a composition with an absorbable gel-forming liquid that transforms to a flexible gel or semi-solid upon contacting water in the biologic environment.

Many drug delivery systems have been developed, tested and utilized for the controlled in vivo release of
30 pharmaceutical compositions. For example, polyesters such as poly(DL-lactic acid), poly(glycolic acid), poly(ϵ -caprolactone) and various other copolymers have been used to release biologically active molecules such as progesterone; these have been in the form of microcapsules, films or rods (M. Chasin and R. Langer, editors, Biodegradable Polymers as Drug Delivery Systems, Dekker, NY
35 1990). Upon implantation of the polymer/therapeutic agent composition, for example, subcutaneously or intramuscularly, the therapeutic agent is released over a

Several EPO publications and U.S. Patents have addressed issues of polymer matrix design and its role in regulating the rate and extent of release of therapeutic agents *in vivo*.

Hutchinson (U.S. Pat. No. 4,767,628) controlled the release of a therapeutic agent by uniform dispersion in a polymeric device. It is disclosed that this formulation provides for controlled continuous release by the overlap
25 of two phases: first, a diffusion-dependent leaching of the drug from the surface of the formulation; and second, releasing by aqueous channels induced by degradation of the polymer.

Other in-situ forming biodegradable implants and
30 methods of forming them are described in U.S. Pat. Nos.
5,278,201 ('201 Patent) and U.S. Pat. No. 5,077,049 ('049
Patent), to Dunn et al. The Dunn et al. patents disclose
methods for assisting the restoration of periodontal tissue
in a periodontal pocket and for retarding a migration of
35 epithelial cells along the root surface of a tooth. The
'049 Patent discloses methods which involve placement of an
in-situ forming biodegradable barrier adjacent to the

surface of the tooth. The barrier is microporous and includes pores of defined size and can include biologically active agents. The barrier formation is achieved by placing a liquid solution of a biodegradable polymer, such as poly(dl-lactide-co-glycolide) water-coagulatable, thermoplastic in a water miscible, non-toxic organic solvent such as N-methyl pyrrolidone (i.e., to achieve a typical polymer concentration of about 50%) into the periodontal pocket. The organic solvent dissipates into the periodontal fluids and the biodegradable, water coagulatable polymer forms an in-situ solid biodegradable implant. The dissipation of solvent creates pores within the solid biodegradable implant to promote cell ingrowth. The '859 Patent likewise discloses methods for the same indications involving the formation of the biodegradable barrier from a liquid mixture of a biodegradable, curable thermosetting prepolymer, curing agent and water-soluble material such as salt, sugar, and water-soluble polymer. The curable thermosetting prepolymer is described as an acrylic-ester terminated absorbable polymer.

In addition, a number of systems for the controlled delivery of biologically active compounds to a variety of sites are disclosed in the literature. For example, U.S. Patent No. 5,011,692, to Fujioka et al., discloses a sustained pulsewise release pharmaceutical preparation which comprises drug-containing polymeric material layers. The polymeric material layers contain the drug only in a slight amount, or free of the drug. The entire surface extends in a direction perpendicular to the layer plane and is coated with a polymeric material which is insoluble in water. These types of pulsewise-release pharmaceutical dosages are suitable for embedding beneath the skin.

U.S. Pat. No. 5,366,756, to Chesterfield et al., describes a method of preparing porous bioabsorbable surgical implant materials. The method comprises providing a quantity of particles of bioabsorbable implant material, and coating particles of bioabsorbable implant material

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with at least one growth factor. The implant can also contain antimicrobial agents.

U.S. Patent No. 5,385,738, to Yamhira et al., discloses a sustained-release injection system, comprising
5 a suspension of a powder comprised of an active ingredient and a pharmaceutically acceptable biodegradable carrier (e.g., proteins, polysaccharides, and synthetic high molecular weight compounds, preferably collagen, atelo collagen, gelatin, and a mixture thereof) in a viscous
10 solvent (e.g., vegetable oils, polyethylene glycol, propylene glycol, silicone oil, and medium-chain fatty acid triglycerides) for injection. The active ingredient in the pharmaceutical formulation is incorporated into the biodegradable carrier in the following state: (i) the
15 active ingredient is chemically bound to the carrier matrix; (ii) the active ingredient is bound to the carrier matrix by intermolecular action; or (iii) the active ingredient is physically embraced within the carrier matrix.

Moreover, such systems as those previously described
20 in the literature, for example, such as by Dunn, et al. (U.S. Pat. No. 4,938,763), teach in-situ formations of biodegradable, microporous, solid implants in a living body through coagulation of a solution of a polymer in an
25 organic solvent such as N-methyl-2-pyrrolidine. However, the use of solvents, including those of low molecular organic ones, facilitates migration of the solution from the application site thereby causing damage to living tissue including cell dehydration and necrosis. Loss of
30 the solvent mass can lead to shrinkage of the coagulum and separation from surrounding tissue.

US Patent No. 5,612,052 describes cation-exchanging microparticles made typically of carboxyl-bearing polyester chains onto which basic bioactive agents are immobilized to
35 provide a control release system within an absorbable gel-forming liquid polyester. The contents of US Patent 5,612,052 is incorporated herein by reference. Conjugating

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SUMMARY OF THE INVENTION

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A preferred bound microparticle of the immediately foregoing, denoted group B, is where said peptide, protein or a combination thereof or a pharmaceutically acceptable salt thereof comprises 0.1% to 30% of the total mass of the bound microparticle.

A preferred bound microparticle of the immediately foregoing, denoted group C, is where said absorbable heterochain polymer core comprises glycolate units.

A preferred bound microparticle of the immediately foregoing, denoted group D, is where the absorbable heterochain polymer core further comprises citrate residues, tartrate residues or malate residues.

A preferred bound microparticle of the immediately foregoing, denoted group E, is where the ratio of glycolate units to citrate residues, to tartrate residues or to malate residues is about 7-1 to about 20-1.

Another preferred bound microparticle of group C is where said glycolate units terminate with a carboxyl moiety.

Yet another preferred bound microparticle of group C is where said glycolate units terminate with an amine moiety.

In another aspect, this invention provides an encased microparticle comprising one or more of a bound microparticle within an absorbable encasing polymer

wherein said bound microparticle comprises an absorbable heterochain polymer core and one or more peptide, one or more protein or a combination thereof immobilized on said absorbable heterochain polymer core,

where each peptide is independently selected from the group consisting of growth hormone releasing peptide (GHRP), luteinizing hormone-releasing hormone (LHRH), somatostatin, bombesin, gastrin releasing peptide (GRP), calcitonin, bradykinin, galanin, melanocyte stimulating hormone (MSH), growth hormone releasing factor (GRF), amylin, tachykinins, secretin, parathyroid hormone (PTH), enkaphelin, endothelin, calcitonin gene releasing peptide

(CGRP), neuromedins, parathyroid hormone related protein (PTHrP), glucagon, neurotensin, adrenocorticotrophic hormone (ACTH), peptide YY (PYY), glucagon releasing peptide (GLP), vasoactive intestinal peptide (VIP),
5 pituitary adenylate cyclase activating peptide (PACAP), motilin, substance P, neuropeptide Y (NPY), TSH and analogs and fragments thereof or a pharmaceutically acceptable salt thereof;

each protein is independently selected from the group
10 consisting of growth hormone, erythropoietin, granulocyte-colony stimulating factor, granulocyte-macrophage-colony stimulating factor and interferons; and where said absorbable heterochain polymer core comprises glycolate units.

15 A preferred encased microparticle of the immediately foregoing is where said peptide, protein or combination thereof or pharmaceutically acceptable salt thereof comprises 0.1% to 30% of the total mass of the bound microparticle, and where said absorbable heterochain
20 polymer core further comprises citrate residues, tartrate residues or malate residues.

A preferred encased microparticle of the immediately foregoing, denoted group F, is where the ratio of glycolate units to citrate residues, to tartrate residues or to
25 malate residues is about 7-1 to about 20-1 and said glycolate units terminate with a carboxyl moiety or an amine moiety.

A preferred encased microparticle of the immediately foregoing is where said absorbable encasing polymer
30 comprises

- (a) l-lactide based units and glycolide based units,
- (b) d,l-lactide based units and glycolide based units,
- (c) d,l-lactide based units or
- (d) l-lactide based units and d,l-lactide based units.

35 A preferred encased microparticle of the immediately foregoing is where the ratio of l-lactide based units to glycolide based units is about 75-25 to about 90-10, the

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ratio of l-lactide based units to d,l-lactide based units is about 80-20 and the ratio of d,l-lactide based units to glycolide based units is about 75-25 to about 90-10.

A preferred encased microparticle of group F is where 5 the absorbable encasing polymer constitutes 5 to 70% of the total mass of the encased microparticle.

A preferred encased microparticle of the immediately foregoing is where the absorbable encasing polymer constitutes 20-60% of the total mass of the encased 10 microparticle.

A preferred encased microparticle of the immediately foregoing is where the absorbable encasing polymer constitutes 30-50% of the total mass of the encased microparticle.

15 In another aspect, this invention provides a pharmaceutical composition comprising the bound microparticles described above and a pharmaceutically acceptable carrier.

In another aspect, this invention provides a 20 pharmaceutical composition comprising the bound microparticles described above, a non-aqueous absorbable gel-forming liquid polyester and optionally a pharmaceutically acceptable carrier.

In another aspect, this invention provides a 25 pharmaceutical composition comprising the encased microparticles described above and a pharmaceutically acceptable carrier.

In another aspect, this invention provides a 30 pharmaceutical composition comprising the encased microparticles described above, a non-aqueous absorbable gel-forming liquid polyester and optionally a pharmaceutically acceptable carrier.

Another preferred bound microparticle of group D, denoted group G, is where the absorbable heterochain 35 polymer core comprises citrate residues and the peptide is an LHRH analog.

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- A preferred bound microparticle of the immediately foregoing is where the ratio of glycolate units to citrate residues of the absorbable heterochain polymer core is about 7-1 to about 20-1 and where the LHRH analog is p-Glu-
5 His-Trp-Ser-Tyr-D-Trp-Leu-Arg-Pro-Gly-NH₂.

Another preferred bound microparticle of group D, denoted group H, is where the absorbable heterochain polymer core comprises tartrate residues and the peptide is an LHRH analog.

- 10 A preferred bound microparticle of the immediately foregoing is where the ratio of glycolate units to tartrate residues is about 7-1 to about 20-1 and the LHRH analog is p-Glu-His-Trp-Ser-Tyr-D-Trp-Leu-Arg-Pro-Gly-NH₂.

- Yet another preferred bound microparticle of group D,
15 denoted group I, is where the absorbable heterochain polymer core comprises citrate residues and the peptide is a somatostatin analog.

- A preferred bound microparticle of the immediately foregoing is where the ratio of glycolate units to citrate
20 residues is about 7-1 to about 20-1 and the somatostatin analog is H- β -D-Nal-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH₂, where the two Cys are bonded by a disulfide bond, N-hydroxyethylpiperazinyl-acetyl-D-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂, where the two Cys are bonded by a disulfide bond
25 or N-hydroxyethylpiperazinyl-ethylsulfonyl-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂, where the two Cys are bonded by a disulfide bond.

- Yet another preferred bound microparticle of group D, denoted group J, is where the absorbable heterochain
30 polymer core comprises tartrate residues and the peptide is a somatostatin analog.

- A preferred bound microparticle of the immediately foregoing is where the ratio of glycolate units to tartrate residues is about 7-1 to about 20-1 and the somatostatin
35 analog is H- β -D-Nal-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH₂, where the two Cys are bonded by a disulfide bond, N-hydroxyethylpiperazinyl-acetyl-D-Phe-Cys-Tyr-D-Trp-Lys-Abu-

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Cys-Thr-NH₂ where the two Cys are bonded by a disulfide bond or N-hydroxyethylpiperazinyl-ethylsulfonyl-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂ where the two Cys are bonded by a disulfide bond.

5 A preferred encased microparticle of this invention is an encased microparticle comprising one or more bound microparticles of group G encased within an absorbable encasing polymer which comprises

- (a) l-lactide based units and glycolide based units,
- 10 (b) d,l-lactide based units and glycolide based units,
- (c) d,l-lactide based units or
- (d) l-lactide based units and d,l-lactide based units.

A preferred encased microparticle of the immediately foregoing is where the ratio of glycolate units to citrate
15 residues of the absorbable polymer core is about 7-1 to about 20-1, the LHRH analog is p-Glu-His-Trp-Ser-Tyr-D-Trp-Leu-Arg-Pro-Gly-NH₂, and where the ratio of:

- (a) l-lactide based units to glycolide based units is about 75-25 to about 90-10,
- 20 (b) d,l-lactide based units to glycolide based units is about 75-25 to about 90-10 and
- (c) l-lactide based units to d,l-lactide based units is about 80-20.

Another preferred encased microparticle comprises one
25 or more bound microparticles of group H encased within an absorbable encasing polymer which comprises

- (a) l-lactide based units and glycolide based units,
- (b) d,l-lactide based units and glycolide based units,
- (c) d,l-lactide based units or
- 30 (d) l-lactide based units and d,l-lactide based units.

A preferred encased microparticle of the immediately foregoing is where the ratio of glycolate units to tartrate residues of the absorbable polymer core is about 7-1 to about 20-1, the LHRH analog is p-Glu-His-Trp-Ser-Tyr-D-Trp-
35 Leu-Arg-Pro-Gly-NH₂, and where the ratio of:

- (a) l-lactide based units to glycolide based units is about 75-25 to about 90-10,

(c) l-lactide based units to d,l-lactide based units is about 80-20.

(a) l-lactide based units and glycolide based units,
(b) d,l-lactide based units and glycolide based units,

(d) l-lactide based units and d,l-lactide based units.

(a) 1-lactide based units to glycolide based units is about 75-25 to about 90-10,

(c) l-lactide based units to d,l-lactide based units is about 80-20.

(a) l-lactide based units and glycolide based units,
(b) d,l-lactide based units and glycolide based units,
(c) d,l-lactide based units or

A preferred encased microparticle of the immediately foregoing is where the ratio of glycolate units to tartrate

35 In yet another aspect, this invention provides a process for making an encased microparticle as described above comprising the step of encasing a bound microparticle

with an absorbable encasing polymer using an emulsion technique.

DETAILED DESCRIPTION

The term "absorbable" as used herein, means a water insoluble material such as a polymer which undergoes chain disassociation in the biological environment to water soluble by-products.

The term "microparticle" as used herein, refers to the particles of absorbable polyester, which are preferably in essentially spherical form.

The term "bound microparticle" as used herein, refers to a microparticle having one or more peptide and/or one or more protein ionically immobilized on the microparticle.

The term "encased microparticle" as used herein, refers to a bound microparticle having a polymer coating, where the polymer coating is not necessarily completely occlusive.

The term "polymer core" as used herein, is another way of referring to microparticles.

The term "encasing polymer" as used herein, refers to the polymer that is used to encase a bound microparticle.

The term "gel-forming liquid polyester" as used herein, refers to materials which absorb solvents such as water, undergo phase transformation and maintain three dimensional networks capable of reversible deformation.

The instant application denotes amino acids using the standard three letter abbreviation known in the art, for example Ala = alanine.

A microparticle of the present invention is crystalline and is made of an absorbable polyester, such as polyglycolide having one or more carboxylic groups on the individual chains which results in a sufficient concentration of carboxylic groups on the surface of the microparticle and immediate subsurface of the microparticle to complex and ionically immobilize a peptide(s) and/or a protein(s) having one or more basic groups. Or the carboxylic groups of the polyglycolide can be amidated, for

example by a diamine, preferably a primary or secondary amine or a mixture thereof, wherein the amine forms a complex that ionically immobilizes a peptide(s) and/or a protein(s) having one or more acidic groups. Since the surface of the microparticles is not necessarily homogeneous, the term "subsurface" refers to the crevices and the like found on the surface of the microparticles. The bound microparticles provide a means for the controlled release of a peptide(s) and/or protein(s) in a patient. To further control the release of the immobilized peptide(s) and/or protein(s), the bound microparticles can be encased individually or in groups with an absorbable polymer coating. The bound microparticles release the peptide(s) and/or protein(s) over a period of about two days to about three months in a patient, preferably about one week to about three months. The encased microparticles release the peptide(s) and/or protein(s) over a period of about three days to six months in a patient, preferably about two weeks to five months.

Typical examples of a peptide that can be immobilized on a microparticle include but are not limited to growth hormone releasing peptide (GHRP), luteinizing hormone-releasing hormone (LHRH), somatostatin, bombesin, gastrin releasing peptide (GRP), calcitonin, bradykinin, galanin, melanocyte stimulating hormone (MSH), growth hormone releasing factor (GRF), amylin, tachykinins, secretin, parathyroid hormone (PTH), enkaphelin, endothelin, calcitonin gene releasing peptide (CGRP), neuromedins, parathyroid hormone related protein (PTHrP), glucagon, neurotensin, adrenocorticotrophic hormone (ACTH), peptide YY (PYY), glucagon releasing peptide (GLP), vasoactive intestinal peptide (VIP), pituitary adenylate cyclase activating peptide (PACAP), motilin, substance P, neuropeptide Y (NPY), TSH, and analogs and fragments thereof. Examples of proteins that can be immobilized on a microparticle are growth hormone, erythropoietin,

granulocyte-colony stimulating factor, granulocyte-macrophage-colony stimulating factor and interferons.

A microparticle can be made of a lactide based polymer or a solid semi-crystalline polylactone such as polyglycolide which can be formed by ring opening polymerization of acid-bearing hydroxylic initiators such as glycolic, lactic, malic, tartaric, and citric acid. A microparticle of the present invention can be synthesized according to the following procedure. In a reaction vessel are mixed a lactide based monomer and/or a lactone such as glycolide and an acid initiator such as tartaric acid, malic acid or citric acid. The reaction vessel is warmed to about 35-45°C, preferably 40°C and put under vacuum for about 20-60 minutes, preferably 30 minutes. The temperature of the reaction vessel is raised to about 105-115°C, preferably 110°C. Once this temperature is reached the vessel is placed under an atmosphere of oxygen-free nitrogen, and the mixture is stirred. Once the mixture melts, a catalytic amount of an organometallic catalyst suitable for ring opening polymerization, such as stannous 2-ethyl-hexanoate solution in a non-protic solvent, such as toluene is added. A vacuum is reapplied for about 30-90 seconds to remove toluene without significant removal of monomer. The temperature of the mixture is raised to about 115-125°C, preferably 120°C for about 5-10 minutes before further raising it to about 145-150°C. It was kept at this temperature for about 3-5 hours, preferably 4 hours, under constant mechanical stirring.

The resulting polymer is micronized by initially grinding it using a Knife-grinder. The polymer is then micronized in an Aljet Micronizer using a pressurized dry nitrogen stream. The mean particle diameter size is analyzed in a Malvern Mastersizer/E using a volume distribution model and 200/5 cS silicone oil as dispersant.

The polymer is purified and the sodium salt thereof is formed by dispersing the micronized polymer in acetone and placing it in a sonicator, preferably for about 30 minutes.

During this time the dispersion was also homogenized at about 8,000-24,000 rpm, preferably 9,500 rpm, using a homogenizer. After this sonication/ homogenization step the dispersion is centrifuged at about 3,000-7,000 rpm, preferably 5,000 rpm _ preferably for about 30 minutes in a centrifuge. The supernatant is discarded, the centrifuge cakes re-suspended in fresh acetone, and the sonication/homogenization step repeated. Once the second centrifugation is complete, the supernatant is discarded and the cakes were re-suspended in deionized water. One final sonication/homogenization step is then carried out to remove any remaining acetone and the dispersion is once again centrifuged at about 5,000 rpm for about 30 minutes.

The centrifuge cakes are re-suspended in fresh deionized water and the pH of the dispersion is monitored. Sufficient volumes of a weak base such as 0.2M sodium carbonate solution are added with stirring to raise the pH to between about pH 8 and about pH 9. The dispersions are allowed to stir for about 30 minutes before being vacuum-filtered over filter paper. The filter cakes are rinsed with further deionized water, frozen, and lyophilized.

Purification is monitored by differential scanning calorimetry (DSC) with a heating rate of about 5°C/min to 15°C/min, preferably 10°C/min.

An anion-exchanger microparticle is obtained by taking the cation-exchanger microparticles and incubating it in hot dilute solution (~80°C) of a diamine, it is preferred that the amines can be both a primary amine or both a secondary amine or a mixture of a primary and a secondary amine, of known concentration in dioxane or THF under an inert gas such as argon. The concentration of the diamine in dioxane or THF is determined by acidimetry. When the reaction practically ceases to take place, the amidated microparticles are separated by filtration, rinsed with dioxane or THF, and dried under reduced pressure.

A peptide(s) and/or protein(s) can be immobilized on a microparticle according to the following method. The

5 filtering out the bound microparticles. The filter cakes are rinsed with further deionized water, frozen, and lyophilized. Samples are then analyzed for nitrogen by elemental analysis to determine the amount of the peptide(s) and/or protein(s) immobilized.

The size of a microparticle plays a role in the amount of a peptide and/or protein that a microparticle of the instant invention can immobilize. The smaller the size of a microparticle, the more surface area a mass of microparticles possess and, thus, the more peptide and/or protein can be immobilized per mass of microparticles. Size reduction of the microparticles to micron or sub-micron dimensions can be achieved as described above. The diameter of the microparticles can range in size from about 0.5 μm to 100 μm , preferably 1 μm to 15 μm and more preferably 3 μm to 10 μm .

The absorbable encasing polymer can be a crystalline or non-crystalline lactide/glycolide copolymer, amorphous l-lactide/d,l-lactide co-polymer, caprolactone/ glycolide copolymer or trimethylene carbonate/glycolide copolymer, 25 that is soluble in conventional organic solvents, such as chloroform, methylene chloride, acetone, acetonitrile, ethyl acetate, and ethyl formate. Non-solvents of such an absorbable encasing polymer include water, low boiling temperature alcohols and hydrocarbons. The absorbable 30 encasing polymers can be synthesized by catalyzing ring-opening polymerization of lactones, or by polymerization of cyclic monomers such as ϵ -caprolactone, p-dioxanone, trimethylene carbonate, 1,5-dioxepan-2-one or 1,4-dioxepan-2-one in the presence of a chain initiator, such as a 35 hydroxy polycarboxylic acid. Still another method involves reacting an organic polycarboxylic acid with a pre-formed

polyester, which is disclosed in U.S. Patent No. 5,612,052, the contents of which is incorporated herein by reference.

The encasing of the bound microparticles can be achieved by phase separation of an emulsion. An alternate
5 encasing method entails the use of an ultrasonic atomizer where a dispersion of the bound microparticles in an absorbable encasing polymer solution is introduced as micro-droplets into a cooled non-solvent medium. Bound microparticles are encased with an absorbable encasing
10 copolymer of lactide and glycolide using traditional microencapsulation or coating techniques of solid particles such as the emulsion evaporation method described by H. Demian and S.W. Shalaby for encapsulating barium sulfate microparticles as disclosed in U.S. Patent application
15 USSN: 08/467,361, the contents of which are incorporated herein by reference, or by coagulation of solid microparticles encased in a polymer solution and delivered through an ultrasonic atomizer (nebulizer) into a liquid medium that is a non-solvent for the encasing polymer, but
20 where the liquid medium non-solvent is capable of extracting the solvent of the encasing polymer solution about the encased solid microparticles. Depending on the concentration of the polymer solution for encasing the microparticles, the number of the original bound
25 microparticles in the encased microparticles can vary from 1 to several hundred with an average diameter of an encased microparticle ranging from 0.5 μm to 100 μm .

The following method relates to the preparation of encased peptide-loaded and/or protein-loaded (hereinafter
30 peptide-loaded) cation exchangers by nebulization. The encasing copolymer of interest is dissolved in a solvent, such as either acetonitrile, ethyl acetate or ethyl formate at a concentration of between 10 and 30% (W/W). A sufficient weight of this solution is used for dispersion
35 of the peptide-loaded CE so that the weight ratio of peptide-loaded CE to encasing copolymer ranges from about 30:70 to about 80:20. Dispersion is achieved by high speed

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homogenization. The dispersion is fed at a flow rate of between 1ml/min and 10 ml/min to an ultrasonic atomization nozzle with variable frequency - this frequency can be altered from 12kHz to 35kHz - higher frequency allows higher flow rates while maintaining particle characteristics. The dispersion is thus nebulized into a collecting sink made up of at least 1 to 10 times excess of isopropanol or ethanol (compared to the volume of encasing copolymer solvent used) containing sufficient dry-ice pellets (usually 0.5 - 1Kg by weight per liter of IPA) so that the temperature of the slurry remains between -70° and -80°C throughout the nebulization. This slurry is stirred at between 300 and 700 rpm depending on its volume. In the case of acetonitrile as solvent, the nebulization droplets will freeze immediately on contact with the slurry. Once nebulization is complete the entire dispersion is allowed to thaw of its own accord to between 10°C and room temperature before vacuum filtering. The filter cakes are rinsed with de-ionized water to remove excess non-solvent. The particles obtained have the appearance of smooth microspheres in the case of a predominantly d,l-lactide encasing copolymer; they appear slightly wrinkled when the encasing copolymer is mainly l-lactide based.

The binding capacity of a microparticle ion-exchanger can be determined as follows. For example, for a cation-exchanger microparticle, available carboxylic groups, in a predetermined mass of the microparticles, are neutralized using cold dilute aqueous sodium carbonate solution of known normality. The neutralized microparticles are isolated by filtration and rinsed thoroughly with cold deionized water and then air dried. The solid microparticles are then incubated in dilute solution of Pilocarpine hydrochloride of known concentration so as to provide a slight excess of the basic drug over that predicted from the binding capacity data. The concentration of the remaining Pilocarpine HCl in the aqueous medium is monitored for a period of time until no

significant change in the base pick-up by the microparticles can be recorded. The percent of immobilized base on the microparticles is determined from the exhaustion data and then verified by elemental analysis for 5 nitrogen.

The binding capacity of the anion-exchanger (amidated particles) is determined by (1) elemental analysis for nitrogen and (2) extent of binding to Naproxen by measuring the extent of Naproxen removed from a dilute solution using HPLC. The latter is confirmed by release of the immobilized Naproxen with a dilute sodium hydroxide solution of known concentration.

The bound microparticles or the encased microparticles of this invention can be administered to a patient via administration routes well known to those of ordinary skill in the art, such as parenteral administration, oral administration or topical administration. Preferably, it is administered as a powder or a suspension via intranasal route or as an inhalant through the pulmonary system. When it is administered parenterally it is preferable that it is administered as a dispersion in an isotonic aqueous medium or in a non-aqueous, absorbable gel-forming liquid polyester as described in U.S. Patent No. 5,612,052, the contents of which are incorporated herein by reference. The formulations comprising bound microparticles and/or encased microparticles of the present invention can also include a variety of optional components. Such components include, but are not limited to, surfactants, viscosity controlling agents, medicinal agents, cell growth modulators, dyes, complexing agents, antioxidants, other polymers such as carboxymethyl cellulose, gums such as guar gum, waxes/oils such as castor oil, glycerol, dibutyl phthalate and di(2-ethylhexyl)phthalate as well as many others. If used, such optional components comprise from about 0.1% to about 20%, preferably from about 0.5% to about 5% of the total formulation.

The effective dosages of bound microparticles or encased microparticles to be administered to a patient can be determined by the attending physician or veterinarian and will be dependent upon the proper dosages contemplated
5 for the peptide(s) and/or protein(s) and the quantity of the peptide(s) and/or protein(s) immobilized on the microparticles. Such dosages will either be known or can be determined by one of ordinary skill in the art.

The preparation of gel-formers is disclosed in US
10 Patent No. 5,612,052, the contents of which is incorporated herein by reference. Specific examples of gel formers are described below.

Preparation of 80/20 (by weight) Block Copolymers of 60/40 Trimethylene

15 Carbonate/Glycolide and Polyethylene Glycol-400 (GF-1): A flame-dried resin kettle equipped with a mechanical stirrer and a nitrogen inlet was charged with polyethylene glycol-400 (0.299 mole, 119.5 g), stannous octoate (0.2 M in toluene, 4.700 ml, 0.946 mmole), glycolide (1.78 mole,
20 206.5 g) and trimethylene carbonate (2.65 mole, 270 g). The reactor was purged with argon several times and then heated to melt and then heated to and stirred at about 150°C for about 12 hours. At the conclusion of the reaction, the temperature was lowered while maintaining
25 fluidity and excess monomer was removed under reduced pressure. The resulting polymer was analyzed by infrared and NMR for composition and gel-permeation chromatography for molecular weight.

Preparation of 15/85 (by weight) Block Copolymer of
30 60/40 Trimethylene

Carbonate/Glycolide and Polyethylene Glycol-400 (GF-2): The title copolymer was synthesized according to the procedure described for GF-1 but using polyethylene glycol-400 (1.063 mole, 425 g), stannous octoate (0.2 M in toluene, 1,760ml,
35 0.35 mmole), glycolide (0.279 mole, 32.4 g) and trimethylene carbonate (0.418 mole, 42.6 g) and stirring for about 9 hours.

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Preparation of 80/20 (by weight) Block Copolymer of 90/10 Trimethylene

Carbonate/Glycolide and Polyethylene Glycol-1500 (GF-3): The title copolymer was synthesized according to the procedure described for GF-1 but using polyethylene glycol-1500 (0.267 mole, 400 g), stannous octoate (0.2 M in toluene, 1200 ml, 0.247 mmole), glycolide (0.097 mole, 11.2 g) and trimethylene carbonate (0.87 mole, 88.7 g) and stirring for about 13 hours.

Example I

Preparation, Micronization, and Purification of Poly(glycolic acid) polymers initiated with Citric Acid (PGCA) for use as Cation Exchangers (CE)

Example I(a): 7/1 PGCA- A 500 ml glass reactor was loaded with 242.63 g of glycolide (Purac Biochem, Arkelsedijk, The Netherlands) and 57.37 g of citric acid (Aldrich, Gillingham, Dorset, U.K.). The citric acid had been further dried over silica gel (Fisher Scientific, Loughborough, Leics., U.K.) in an Abderhalden apparatus (Aldrich, St. Louis, Missouri, USA). The reactor was immersed in an oil bath at about 40°C and put under vacuum (0.04 mbar) for about 30 minutes. The bath was then lowered and it's temperature raised to about 110°C. Once this temperature was reached the reactor was placed under an atmosphere of oxygen-free nitrogen and re-immersed. The contents were stirred at about 100 rpm using a Heidolph stirrer (Heidolph Elektro GmbH, Kelheim, Germany). Once the reactor contents melted 1.09 ml of a 0.1M stannous 2-ethyl-hexanoate solution (Sigma, St. Louis, Missouri, USA) in toluene (Riedel de-Haen, Seelze, Germany) was added (stoichiometric ratio of 50 ppm). A vacuum was reapplied via a liquid nitrogen trap for about 30 seconds to remove toluene without significant removal of monomer. The oil bath temperature was then raised to about 120°C for about 5 minutes before further raising it to about 150°C. It was kept at this temperature for about 4 hours under constant

mechanical stirring of about 100 rpm. The title polymer was obtained.

Example I(b): 10/1 PGCA- The title polymer was obtained by following the procedure of Example Ia, but using 257.40 g of glycolide, 42.60 g of citric acid and 1.10 ml of a 0.1M stannous 2-ethyl-hexanoate solution in toluene (stoichiometric ratio of 50 ppm).

Example I(c): 15/1 PGCA- 15/1 PGCA- A flame-dried resin kettle equipped with a mechanical stirrer and an argon inlet was charged with glycolide (2.586 mole, 300 g), anhydrous citric acid (0.172 mole, 33 g), and stannous octoate (0.2 M in toluene, 862ml, 0.172 mmole). The polymerization reactor and its contents were purged with dry argon several times. After melting the polymerization charge, the reactants were heated and stirred at about 160°C until the polymer started to precipitate from the melt. Shortly after partial precipitation, the stirring was terminated and the reaction was continued at about 160°C for about 2 hours. At the conclusion of the polymerization, the temperature was lowered below 120°C and excess monomer was removed under reduced pressure. The composition of the isolated polymer was verified using infrared and NMR spectroscopy.

Micronization- Each of the polymers of Examples I(a), I(b) and I(c) were ground initially using a Knife-grinder (IKA, Staufen, Germany). They were then micronized in an Aljet Micronizer (Fluid Energy Aljet, Plumsteadville, Pennsylvania, USA) using a pressurized dry nitrogen stream. Example I(a) had a mean particle diameter size of 24.84 μm by analysis in a Malvern Mastersizer/E (Malvern, Worcs., U.K.) using a volume distribution model and 200/5 cS silicone oil (Dow Corning, Seneffe, Belgium) as dispersant. Examples I(b) and I(c) had mean particle diameter sizes of 4.69 μm and 6.31 μm , respectively, after micronization.

Purification/Sodium Salt Formation- Fifty gram batches of Examples I(a), I(b), and I(c) were dispersed in 2L of acetone (Riedel de-Haen, Seelze, Germany) and placed in a

The centrifuge cakes were re-suspended in fresh deionized water and the pH of the dispersion was monitored. Sufficient volumes of 0.2M sodium carbonate solution were added in each case (with stirring) to raise the pH to 20 between about pH 8 and about pH 9. The dispersions were allowed to stir for about 30 minutes before being vacuum-filtered over a Whatman no.1 (24 cm diameter) filter paper (Whatman Intl. Ltd., Maidstone, Kent, U.K.). The filter cakes were rinsed with further deionized water, frozen, and 25 lyophilized in an Edwards SuperModulyo Lyophilizer (Edwards, Crawley, West Sussex, U.K.).

Purification was monitored by differential scanning calorimetry (DSC) using a TA DSC912S (TA Instruments, New Castle, Delaware, USA) with a heating rate of 10°C/min. The DSC thermograms obtained in each case did not show any endothermic peak for monomeric glycolide but showed endotherms at 176°C, 178°C, and 180°C for Examples I(a), I(b), and I(c), respectively.

Example II

**Preparation of Microparticulate Cation-Exchanger of
Glycolide/Malic Acid Copolymer PGMA**

- The title microparticle was synthesized according to the method described in Example I(c) but using glycolide (2.586 mole, 300 g), anhydrous malic acid (0.172 mole, 23 g), and stannous octoate (0.2 M in toluene, 862ml, 0.172 m mole). Differential Scanning Calorimetry was used to determine the polymer melting temperature ($T_m = 206^\circ\text{C}$).
- The solid polymer was ground to achieve average particle diameter of about 125 μm using a Wiley mill. Further reduction of the particle size to about 5-10 μm diameter was achieved using a jet-mill receiving pressurized dry nitrogen. The resulting microparticles were rinsed with acetone to remove trace monomer and low molecular weight oligomers. The product was then dried under reduced pressure at 40°C until used. The average diameter of the dry microparticle was determined using a particle size analyzer.

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Example III

**Preparation, Micronization, and Purification of a
Poly(glycolic acid) polymer initiated with Tartaric Acid
(PGTA) for use as a Cation Exchanger (CE)**

- Example III(a): 10/1 PGTA-** A 500 ml glass reactor was loaded with 264.65 g of glycolide (Purac Biochem, Arkelsedijk, The Netherlands) and 34.22 g of L-Tartaric acid (Riedel de-Haen, Seelze, Germany). The tartaric acid had been further dried over silica gel (Fisher Scientific, Loughborough, Leics., U.K.) in an Abderhalden apparatus (Aldrich, St. Louis, MO). The reactor was immersed in an oil bath at about 40°C and put under vacuum (0.04 mbar) for about 30 minutes. The bath was then lowered and it's temperature raised to about 110°C . Once this temperature was reached the reactor was placed under an atmosphere of oxygen-free nitrogen and re-immersed. The contents were stirred at about 100 rpm using a Heidolph stirrer (Heidolph Elektro GmbH, Kelheim, Germany). Once the reactor contents

melted 1.14 ml of a 0.1M stannous 2-ethyl-hexanoate solution (Sigma, St. Louis, Missouri, USA) in toluene (Riedel de-Haen, Seelze, Germany) was added (stoichiometric ratio of 50 ppm). A vacuum was reapplied via a liquid nitrogen trap for about 30 seconds to remove toluene without significant removal of monomer. The oil bath temperature was then raised to about 120°C for about 5 minutes before further raising it to about 150°C. It was kept at this temperature for about 4 hours under constant mechanical stirring of about 100 rpm. The title polymer was obtained.

Micronization—Example III(a) was ground initially using a Knife-grinder (IKA, Staufen, Germany). It was then micronized in an Aljet Micronizer (Fluid Energy Aljet, Plumsteadville, Pennsylvania, USA) using a pressurized dry nitrogen stream. This gave a mean particle diameter of 12.42 μm by analysis in a Malvern Mastersizer/E (Malvern, Worcs., U.K.) using a volume distribution model and 200/5 cS silicone oil (Dow Corning, Seneffe, Belgium) as dispersant.

Purification/Sodium Salt Formation—A 50 g batch of Example III(a) was dispersed in 2L of acetone (Riedel de-Haen) and placed in a sonicator (Branson Ultrasonics BV, Soest, The Netherlands) for about 30 minutes. During this time the dispersion was also homogenized at about 9,500 rpm using an Ultra-turrax T25 homogenizer (IKA, Staufen, Germany). After this sonication/homogenization step the dispersion was centrifuged at about 5,000 rpm for about 30 minutes in a Sorvall centrifuge (Sorvall, Wilmington, Delaware, USA). The supernatant was discarded, the centrifuge cakes re-suspended in fresh acetone, and the sonication/homogenization step repeated. Once the second centrifugation was complete, the supernatant was discarded and the cakes were re-suspended in deionized water. One final sonication/homogenization step was then carried out to remove any remaining acetone and the dispersion was once again centrifuged at about 5,000 rpm for about 30 minutes.

The centrifuge cakes were resuspended in fresh de-ionized water and the pH of the dispersion was monitored. A sufficient volume of 0.2M sodium carbonate solution was added to raise the pH to between about pH 8 and about pH 9.

5 The dispersion was allowed to stir for about 30 minutes before being vacuum-filtered over a Whatman no.1 (24 cm diameter) filter paper (Whatman Intl. Ltd., Maidstone, Kent, U.K.). The filter cake was rinsed with further deionized water, frozen, and lyophilized in an Edwards

10 SuperModulyo Lyophilizer (Edwards, Crawley, West Sussex, U.K.).

Purification was monitored by DSC using a TA DSC912S (TA Instruments New Castle, Delaware, USA) with a heating rate of about 10°C/min. The DSC thermogram obtained did not

15 show any endothermic peak for monomeric glycolide but showed an endotherm at 181°C.

Example III(b): 15/1 PGTA- The title polymer was synthesized according to the procedure described for Example I(c) but using glycolide (2.586 mole, 300 g),

20 anhydrous tartaric acid (0.172 mole, 26.8 g) and stannous octoate (0.2 M in toluene, 862 ml, .0172 mmole). Differential Scanning Calorimetry was used to determine the polymer melting temperature ($T_m = 204^\circ\text{C}$).

The solid polymer was ground to achieve average

25 particle diameter of about 125 μm using a Wiley mill. Further reduction of the particle size to about 5-10 μm diameter was achieved using a jet-mill receiving pressurized dry nitrogen. The resulting microparticles were rinsed with acetone to remove trace amounts of monomer

30 and low molecular weight oligomers. The product was then dried under reduced pressure at about 40°C until used. The average diameter of the dry microparticle was determined using a particle size analyzer.

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Example IV**Preparation of Polyglycolide-based Microparticulate
Anion-Exchanger (AE-1)**

The preparation of an anion-exchanger is achieved in two steps. First, low molecular weight polyglycolide is prepared using a similar procedure in Example I(c), but using the following polymerization charge: glycolide (1 mole, 116 g), 1,3 propanediol as an initiator (30 mmole, 2.22 g) and stannous octoate (0.03 mmole). The size reduction and purification of the polymer are then conducted as also described in Example I(c). In the second step, the practically non-ionic microparticles are incubated in hot dilute solution (~80°C) of a diamine, for example hexanediamine of known concentration in dioxane under argon. The concentration of the diamine in dioxane is determined by acidimetry. When the reaction practically ceases to take place, the amidated microparticles are separated by filtration, rinsed with dioxane, and dried under reduced pressure. The binding capacity of the anion-exchanger (amidated particles) is determined by (1) elemental analysis for nitrogen and (2) extent of binding to Naproxen by measuring the extent of drug removed from a dilute solution using HPLC. The latter is confirmed by release of the immobilized Naproxen with a dilute sodium hydroxide solution of known concentration.

Example V**Preparation of Poly(lactide co-glycolide) copolymers
initiated with propanediol (PLGPD) for use as encasing
materials**

Example V(a): 75/25 P(1)LGPD- A 500 ml glass reactor was loaded with 235.01 g of 1-lactide (Purac Biochem, Arkelsedijk, The Netherlands), 63.09 g of glycolide (Purac Biochem, Arkelsedijk, The Netherlands) and 1.90 g of propanediol (Riedel de-haen, Seelze, Germany) and then 3.96 ml of a 0.1M stannous 2-ethyl-hexanoate solution (Sigma, St. Louis, Missouri, USA) in toluene (Riedel de-haen, Seelze, Germany) was added (stoichiometric ratio of 200

ppm). After drying under vacuum for about one hour to remove the toluene, the reactor was placed under an atmosphere of oxygen-free nitrogen and immersed in an oil bath preheated at about 160°C. The reactor contents were stirred at about 100 rpm with a Heidolph stirrer (Heidolph Elektro GmbH, Kelheim, Germany). Once the contents had melted the temperature was increased to about 180°C and maintained at this level for about 3 hours. An amorphous copolymer was obtained. The copolymer was found to have a molecular weight (MW) of about 12,500 g/mol by gel permeation chromatography (GPC) on a Waters 510 Pump, Waters 410 Differential Refractometer (Waters, Milford, Massachusetts, USA) with light-scattering detection on a Wyatt Minidawn Light Scattering Detector (Wyatt Technology Corporation, Santa Barbara, California, USA).

Example V(b): 90/10 P(1)LGPD- The title product was synthesized according to the procedure of Example V(a) but using 274.31 g of l-lactide, 24.55 g of glycolide, 1.14 g of propanediol and 3.89 ml of a 0.1M stannous 2-ethyl-hexanoate solution in toluene (stoichiometric ratio of 200 ppm). A crystalline copolymer was obtained. The copolymer was found to have a molecular weight of about 20,780 g/mol by GPC.

Example V(c): 90/10 P(d,l)LGPD- The title product was obtained by following the procedure of Example V(a) but using 274.31 g of d,l-lactide, 24.55 g of glycolide, 1.14 g of propanediol and 3.86 ml of a 0.1M stannous 2-ethyl-hexanoate solution in toluene (stoichiometric ratio of 200 ppm). An amorphous copolymer was obtained. The copolymer was found to have a molecular weight of about 20,650 g/mol by GPC.

Example V(d): Poly(l-lactide co-d,l-lactide) copolymer initiated with propanediol (PLGPD) for use as coating material, 80/20 P(1)L(d,l)LPD

The title product was obtained by following the procedure of Example V(a) but using 239.09g of l-lactide, 59.77g of d,l-lactide (Purac Biochem, Arkelsedijk, The

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Netherlands) and 1.14g of propanediol and 3.96 ml of a 0.1M stannous 2-ethyl-hexanoate solution in toluene was added (stoichiometric ratio of 200ppm). An amorphous copolymer was obtained. The copolymer was found to have a molecular weight (Mw) of 22,320 g/mol by GPC. It showed a glassy transition at 48°C by DSC.

Purification— Examples V(a), V(b), and V(c) were each washed by nebulization of a 30% (W/W) solution in acetonitrile (Labscan, Dublin, Ireland) at 8 ml/min into deionized water cooled to about 2°C in a 6L jacketed reactor linked to a circulation bath and stirred at about 350 rpm with a Heidolph stirrer (Heidolph Elektro GmbH, Kelheim, Germany). The solutions were fed to a Vibra-Cell VC 50 Atomization nozzle (Bioblock, Illkirch, France) using a Masterflex pump (Cole Parmer Instrument Co., Niles, Illinois, USA) and nebulization was achieved using a sonication frequency of 12 kHz. The dispersions obtained were filtered over Whatman No.1 (24 cm diameter) filter papers (Whatman Intl. Ltd., Maidstone, Kent, U.K.) and the filter cakes were rinsed with deionized water, frozen, and lyophilized in an Edwards SuperModulyo Lyophilizer (Edwards, Crawley, West Sussex, U.K.).

Purity was confirmed by DSC using a TA DSC912s (TA Instruments, New Castle, Delaware, USA) with a heating rate of 10°C/min which showed glass transitions (Tg) at 44°C, 49°C, 45°C and 48°C for Examples V(a), V(b), V(c) and V(d), respectively.

Example VI

Preparation of Peptide-Loaded Cation Exchangers

Example VI(a): Loading with Peptide A (p-Glu-His-Trp-Ser-Tyr-D-Trp-Leu-Arg-Pro-Gly-NH₂, an LHRH analog)— Four grams of each of the sodium salts of Examples I(a), I(b), I(c) and II(a) were dispersed in solutions containing 1.33 g of the free-base of Peptide A (Kinerton Ltd., Dublin, Ireland) dissolved in 70 ml of deionized water. The dispersions were incubated at room temperature with stirring for about 2 hours before filtering over a 9 cm diameter Whatman No.1

filter paper (Whatman Intl. Ltd., Maidstone, Kent, U.K.). The filter cakes were rinsed with further deionized water, frozen, and lyophilized in an Edwards SuperModulyo (Edwards, Crawley, West Sussex, U.K.). Samples were then analyzed for nitrogen by elemental analysis to determine the amount of Peptide A bound. The following results were obtained :

Examp e	CE Ex.-#	CE Polymer	wt. % Peptide A Bound
10 VI(a)(i)	I(a)	7/1 PGCA	24.52%
VI(a)(ii)	I(b)	10/1 PGCA	12.60%
15 VI(a)(iii)	I(c)	15/1 PGCA	19.29%
VI(a)(iv)	III(a)	10/1 PGTA	17.60%

- 20 **Example VI(b) : Loading with Peptide B** (H- β -D-Nal-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH₂, the two Cys are bonded by a disulfide bond, a somatostatin analogue)- Following the procedure of Example VI(b) and using 4 g of each of the sodium salts of Examples I(a), I(b), I(c) and II(a) and
- 25 1.33 g of the free-base of Peptide B (Kinerton Ltd., Dublin, Ireland) bound microparticles of Examples I(a), I(b) and I(c) with peptide B immobilized thereon were obtained. Samples were analyzed for nitrogen content by elemental analysis to determine the amount of Peptide B

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bound. The results obtained are shown below :

Examp le	CE Ex.-#	CE Polymer	wt. % Peptide B Bound
5 VI(b) (i)	I(a)	7/1 PGCA	25.20%
VI(b) (ii)	I(b)	10/1 PGCA	13.10%
VI(b) (iii)	I(c)	15/1 PGCA	19.64%
10 VI(b) (iv)	III(a)	10/1 PGTA	14.23%

Example VII

Preparation of Encased Polypeptide-Loaded Cation Exchangers by Nebulization

- 15 Polypeptide-loaded cation exchangers were dispersed in acetonitrile (Labscan, Dublin, Ireland) solutions of encasing copolymers, indicated below. This dispersal was achieved by homogenizing with an Ultra-turrax T25 (IKA, Staufen, Germany) at about 9,500 rpm for about 5 minutes. The concentration of the encasing copolymer/ acetonitrile solutions ranged from 12.5% to 25% (W/W) and the ratio of encasing copolymer to polypeptide-loaded cation exchanger ranged from 1:1 to 1.3:1 by weight.
- 20
- 25 After dispersal, the dispersion was fed to a Vibra-Cell VC50 atomization nozzle (Bioblock, Illkirch, France) with a sonication frequency of 16 kHz using a ceramic piston pump (FMI, Oyster Bay, N.Y., USA) set at 2ml/min flow rate. Upon reaching the nozzle the dispersion was
- 30 nebulized into isopropyl alcohol (IPA) (Labscan, Dublin, Ireland) cooled to about -80°C by the addition of dry-ice pellets (A.I.G., Dublin, Ireland). The IPA acted as a collecting non-solvent and was stirred at about 300 rpm using a Heidolph stirrer (Heidolph Elektro GmbH, Kelheim, Germany). Once nebulization was complete the entire dispersion was allowed to thaw to a temperature between about 10°C and about room temperature. The encased
- 35

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microparticles were then recovered by vacuum filtration over a Whatman No.1 filter paper (Whatman Intl. Ltd., Maidstone, Kent, U.K.). The filter cake was rinsed with deionized water, frozen and lyophilized in an Edwards
5 SuperModulyo lyophilizer (Edwards, Crawley, West Sussex, U.K.). The resulting encased microparticles were analyzed for size using the Malvern Mastersizer/E (Malvern, Worcs., U.K.) and 1% Tween 20 in water as a dispersant. The encased
10 microparticles were also analyzed for nitrogen content by elemental analysis to determine peptide content.

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The table below represents the various encasing experiments carried out :

	Ex. #	Peptide-loaded CE: Ex-#	Encasing Copolymer Ex-#	Conc. (W/W) of Encasing Copolymer in Acetonitrile	Encasing Copolymer : Peptide-loaded CE	Mean Particle Diameter	wt.% Peptide Loading
5	VII(a)	VI(a)(ii)	V(a)	24.31%	1:1	122.14 μ m	5.38% Peptide A
	VII(b)	VI(a)(ii)	V(b)	22.41%	1:1	120.15 μ m	6.38% Peptide A
	VII(c)	VI(a)(iii)	V(b)	12.5%	1:1	79.30 μ m	7.76% Peptide A
10	VII(d)	VI(a)(iii)	V(c)	12.5%	1:1	77.85 μ m	8.93% Peptide A
	VII(e)	VI(a)(iv)	V(c)	14.95%	1:1	136.74 μ m	8.75% Peptide A
15	VII(f)	VI(a)(i)	V(c)	14.92%	1.27:1	80.59 μ m	10.31% Peptide A
	VII(g)	VI(b)(ii)	V(a)	25.37%	1:1	140.58 μ m	2.63% Peptide B
	VII(h)	VI(b)(ii)	V(b)	20%	1.15:1	96.77 μ m	5.98% Peptide B
20	VII(i)	VI(b)(iii)	V(b)	12.5%	1:1	102.56 μ m	7.69% Peptide B
	VII(j)	VI(b)(iii)	V(c)	12.5%	1:1	83.72 μ m	7.90% Peptide B
25	VII(k)	VI(b)(iv)	V(c)	14.95%	1:1	135.14 μ m	6.69% Peptide B
	VII(l)	VI(b)(i)	V(c)	14.92%	1.26:1	123.18 μ m	10.11% Peptide B

All samples were sieved over a 180 μ m sieve (Bioblock, Illkirch, France) prior to *in vivo* and/or *in vitro* testing.

A bound microparticle or encased microparticle can be tested *in vitro* to assess the release rate of a bound peptide or bound protein by the following method. An

aliquot of a bound microparticle or encased microparticle having a mass of about 50 mg is placed in a continuous flow-cell system where a buffered phosphate solution at about pH 7.2 and at about 37°C flow across the entire mass of the bound microparticles or encased microparticles at a rate of about 45 ml/hr. Samples of the buffer containing the released drug are collected at about 4°C and analyzed for the peptide or protein concentrations at 1- or 2-day intervals. The release profile of each microparticle is determined over a period of 2 weeks.

A bound microparticle or encased microparticle can be tested to assess the release rate of a bound peptide or bound protein in an *in vivo* system by the following method. Samples are administered to male Wistar rats (Bioresources, 15 Trinity College, Dublin, Ireland) by intramuscular injection to the thigh. The suspension medium consists of 3% carboxymethylcellulose and 1% Tween 20 in saline solution. For Peptide A-loaded samples the effective equivalent dose is 40 $\mu\text{g/Kg/day}$. The dose for Peptide B- 20 loaded samples is 1 mg/Kg/day. Samples are taken by cardiac puncture and the plasma peptide levels are monitored by radioimmunoassays (RIA) specific for Peptide A and Peptide B. In the case of Peptide A-loaded samples (Peptide A is an LHRH analog), a testosterone RIA is also used to monitor 25 testosterone suppression. As an alternative to the suspension medium, gel-formers can be used in certain

cases. The results are shown in Tables A and B, below.

Table A

	Peptide A Examples	Peptide A (>150 pg/ml) Days	Testosterone (<1ng/ml) Days
5	VII(a)	20	21
	VII(b)	10	10
	VII(c)	2	11
	VII(d)	2	11
	VII(e)	2	13
10	VII(f)	2	16
	VII(a) in gel- former	25	44

Table B

	Peptide B Examples	Peptide B (>1000 pg/ml) Days
15	VII(g)	Not tested
	VII(h)	Not tested
	VII(i)	Not tested
20	VII(j)	15
	VII(k)	10
	VII(l)	10

Example VIII

- 25 **Example VIII(a): Nebulization using acetonitrile as solvent and room temperature IPA as non-solvent**

About 1.06 g of the cation exchanger of Example I(c) (not bound to polypeptide) was dispersed in a 25.24% (W/W) solution of encasing copolymer of Example V(a) in
 30 acetonitrile (Labscan, Dublin, Ireland) such that the ratio of cation exchanger to encasing copolymer was about 1.03:1 by weight. This dispersal was achieved by homogenizing with an Ultra-turrax T25 (IKA, Staufen, Germany) at about 9,500rpm for about 5 minutes.

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The nebulization was carried out substantially according to the procedure of Example VIII(a) but using about 0.99g of cation exchanger of Example I(c) (not bound to polypeptide) dispersed in a 24.88% (W/W) solution of encasing copolymer of Example V(a) in ethyl acetate (Riedel-de Haen, Seelze, Germany) such that the ratio of cation exchanger to encasing copolymer was about 0.96:1 by weight. The resulting particles had a mean particle size (d(0.5)) of 100.56 μ m.

35 About 1.02g of cation exchanger of Example I(c) (not bound to polypeptide) was dispersed in a 15.14% (W/W) solution of encasing copolymer of Example V(a) in ethyl

acetate (Riedel-de Haen) such that the ratio of cation exchanger to encasing copolymer was about 1.05:1 by weight. This dispersal was achieved by homogenizing with an Ultraturrax T25 (IKA, Staufen, Germany) at about 9,500 rpm for about 5 minutes.

After dispersal, the dispersion was fed to a Martin
Walter 400 GSIP nebulizer (Sodeva, Le Bouget du Lac,
France) with an ultrasonic frequency setting of about
34.6kHz using a ceramic piston pump (FMI, Oyster Bay, N.Y.
U.S.A.) set at 5ml/min flow rate. Upon reaching the nozzle
the dispersion was nebulized into IPA (Labscan, Dublin,
Ireland) cooled to about -77° by the addition of dry-ice
pellets (A.I.G., Dublin, Ireland). This IPA acted as a
collecting non-solvent and was stirred at 300rpm using a
Heidolph stirrer (Heidolph Elektro GmbH, Kelheim, Germany)
Once nebulization was complete the coated particles were
recovered by vacuum filtration over a Whatman No. 1 filter
paper (Whatman Intl. Ltd., Maidstone, Kent, U.K.). The
filter cake was rinsed with deionized water, frozen and
lyophilized in an Edwards SuperModulyo lyophilizer
(Edwards, Crawley, West Sussex, U.K.). The resulting
particles were analyzed for particle size using the Malvern
Masterizer/E (Malvern, Worcs., U.K.) and 1% Tween 20 in
water as a dispersant. The resulting particles had a mean
particle size (d(0.5)) of 95.69µm.

Example IX

Binding to cation exchanger and subsequent encasing of Somatostatin Analog Peptides C and D

Example IX(a): Loading with peptide C

- 5 About 1.01g of the sodium salt of Example I(c) dispersed in a solution containing 0.25g of the free base of peptide C, which has the structure N-hydroxyethylpiperazinyl-acetyl-D-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂, where the two Cys residues are bonded by a
- 10 disulfide bond (Kinerton Ltd., Dublin, Ireland), dissolved in 40 ml deionized water. The dispersion was incubated with stirring for about 2 hours before filtering over a 9 cm diameter Whatman No. 1 filter paper (Whatman Intl. Ltd., Maidstone, Kent, U.K.). The filter cake was rinsed with
- 15 further deionized water, frozen, and lyophilized in an Edwards SuperModulyo (Edwards, Crawley, West Sussex. U.K.). The sample was then sent for nitrogen analysis to determine the amount of peptide bound, 20.21%.

Example IX(b): Loading with peptide D

- 20 Using the procedure of Example IX(a) but using about 2.04 g of the sodium salt of Example I(c) dispersed in a solution containing 0.51g of the free base of peptide D, which has the structure N-hydroxyethylpiperazinyl-ethylsulfonyl-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂, where
- 25 the two Cys residues are bonded by a disulfide bond (Kinerton Ltd., Dublin, Ireland), dissolved in 80 ml deionized water. The sample was then sent for nitrogen analysis to determine the amount of peptide bound, 19.53%.

Example X

- 30 The bound microparticles of Examples IX(a) and IX(b) were encased as described in Example VII yielding the following results:

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Ex . No.	Peptide- loaded CE	Coating copolymer	Conc. (W/W) of coating copolymer in acetonitrile	Coating copolymer:Peptide loaded CE	Mean Part icle Size (μm)	Wt. % Peptide Loadi ng
X(a)	IX(a)	V(c)	12.51%	1:1	83.33	9.48%
X(b)	IX(b)	V(c)	12.48%	0.98:1	72.15	8.87%
5 X(c)	IX(b)	V(d)	12.35%	0.98:1	86.03	6.74%

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Example XI

Preparation of a Gel-Former Formulation

Encased microparticles of Example VII(a) (0.3 g) were mixed with a liquid gel former (2.0 mL of a 50/50 mixture of component "A" of Example I and component C of Example III, both of which are disclosed in U.S. Patent No. 5,612,052) in a 5 mL syringe barrel using a mechanical micromixer at about 20 rpm for about 10 minutes. The gel-former was presterilized by dry heat and the mixing was conducted using a sterilized stirrer in a laminar flow hood. The formulation was extruded from the 5 mL syringe (after introducing the plunger) into a smaller syringe which is intended for use in administering the formulation. The uniformity of the formulation was checked using optical microscopy. The small syringes were assembled for storage in a dry package and kept at about 4°C until use.

CLAIMS

What is claimed is:

1. A bound microparticle comprising an absorbable
5 heterochain polymer core and one or more peptide, one or more protein or a combination thereof immobilized on said absorbable heterochain polymer core,

wherein each peptide is independently selected from the group consisting of growth hormone releasing peptide
10 (GHRP), luteinizing hormone-releasing hormone (LHRH), somatostatin, bombesin, gastrin releasing peptide (GRP), calcitonin, bradykinin, galanin, melanocyte stimulating hormone (MSH), growth hormone releasing factor (GRF), amylin, tachykinins, secretin, parathyroid hormone (PTH),
15 enkaphelin, endothelin, calcitonin gene releasing peptide (CGRP), neuromedins, parathyroid hormone related protein (PTHrP), glucagon, neurotensin, adrenocorticotrophic hormone (ACTH), peptide YY (PYY), glucagon releasing peptide (GLP), vasoactive intestinal peptide (VIP),
20 pituitary adenylate cyclase activating peptide (PACAP), motilin, substance P, neuropeptide Y (NPY), TSH and analogs and fragments thereof or a pharmaceutically acceptable salt thereof; and

wherein each protein is independently selected from
25 the group consisting of growth hormone, erythropoietin, granulocyte-colony stimulating factor, granulocyte-macrophage-colony stimulating factor and interferons.

2. A bound microparticle according to claim 1 wherein said peptide, protein or a combination thereof or
30 a pharmaceutically acceptable salt thereof comprises 0.1% to 30% of the total mass of the bound microparticle.

3. A bound microparticle according to claim 2 wherein said absorbable heterochain polymer core comprises glycolate units.

- 35 4. A bound microparticle according to claim 3 wherein the absorbable heterochain polymer core further comprises citrate residues.

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5. A bound microparticle according to claim 4 wherein the ratio of glycolate units to citrate residues is about 7-1 to about 20-1.

6. A bound microparticle according to claim 3 wherein the absorbable polymer core further comprises tartrate residues.

7. A bound microparticle according to claim 6 wherein the ratio of glycolate units to tartrate residues is about 7-1 to about 20-1.

8. A bound microparticle according to claim 3 wherein the absorbable heterochain polymer core further comprises malate residues.

9. A bound microparticle according to claim 8 wherein the ratio of glycolate units to malate residues is about 7-1 to about 20-1.

10. A bound microparticle according to claim 3 wherein said glycolate units terminate with a carboxyl moiety.

11. A bound microparticle according to claim 3 wherein said glycolate units terminate with an amine moiety.

12. An encased microparticle comprising one or more of a bound microparticle encased within an absorbable encasing polymer,

wherein said bound microparticle comprises an absorbable heterochain polymer core and one or more peptide, one or more protein or a combination thereof immobilized on said absorbable heterochain polymer core,

where each peptide is independently selected from the group consisting of growth hormone releasing peptide (GHRP), luteinizing hormone-releasing hormone (LHRH), somatostatin, bombesin, gastrin releasing peptide (GRP), calcitonin, bradykinin, galanin, melanocyte stimulating hormone (MSH), growth hormone releasing factor (GRF), amylin, tachykinins, secretin, parathyroid hormone (PTH), enkaphelin, endothelin, calcitonin gene releasing peptide (CGRP), neuromedins, parathyroid hormone related protein

(PTHrP), glucagon, neurotensin, adrenocorticotrophic hormone (ACTH), peptide YY (PYY), glucagon releasing peptide (GLP), vasoactive intestinal peptide (VIP), pituitary adenylate cyclase activating peptide (PACAP),
5 motilin, substance P, neuropeptide Y (NPY), TSH and analogs and fragments thereof or a pharmaceutically acceptable salt thereof;

where each protein is independently selected from the group consisting of growth hormone, erythropoietin,
10 granulocyte-colony stimulating factor, granulocyte-macrophage-colony stimulating factor and interferons; and where said absorbable heterochain polymer core comprises glycolate units.

13. An encased microparticle according to claim 12 wherein said peptide, protein or combination thereof or pharmaceutically acceptable salt thereof comprises 0.1% to 30% of the total mass of the bound microparticle, and where said absorbable heterochain polymer core further comprises citrate residues, tartrate residues or malate residues.

20 14. An encased microparticle according to claim 13 wherein the ratio of glycolate units to citrate residues, tartrate residues or malate residues is about 7-1 to about 20-1 and said glycolate units terminate with a carboxyl moiety or an amine moiety.

25 15. An encased microparticle according to claim 14 wherein said absorbable encasing polymer comprises

- (a) l-lactide based units and glycolide based units,
- (b) d,l-lactide based units and glycolide based units,
- (c) d,l-lactide based units or
- 30 (d) l-lactide based units and d,l-lactide based units.

16. An encased microparticle according to claim 15 wherein the ratio of l-lactide based units to glycolide based units is about 75-25 to about 90-10.

17. An encased microparticle according to claim 15
35 wherein the ratio of l-lactide based units to d,l-lactide based units is about 80-20.

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18. An encased microparticle according to claim 15 wherein the ratio of d,l-lactide based units to glycolide based units is about 75-25 to about 90-10.

19. An encased microparticle according to claim 14 wherein the absorbable encasing polymer constitutes 5 to 70% of the total mass of the encased microparticle.

20. An encased microparticle according to claim 19 wherein the absorbable encasing polymer constitutes 20-60% of the total mass of the encased microparticle.

21. An encased microparticle according to claim 20 wherein the absorbable encasing polymer constitutes 30-50% of the total mass of the encased microparticle.

22. A pharmaceutical composition comprising the bound microparticles according to claim 1 and a pharmaceutically acceptable carrier.

23. A pharmaceutical composition comprising the bound microparticles according to claim 1, a non-aqueous absorbable gel-forming liquid polyester and optionally a pharmaceutically acceptable carrier.

24. A pharmaceutical composition comprising the encased microparticles according to claim 12 and a pharmaceutically acceptable carrier.

25. A pharmaceutical composition comprising the encased microparticles according to claim 12, a non-aqueous absorbable gel-forming liquid polyester and optionally a pharmaceutically acceptable carrier.

26. A bound microparticle according to claim 4 wherein the peptide is an LHRH analog.

27. A bound microparticle according to claim 26 wherein the ratio of glycolate units to citrate residues of the absorbable heterochain polymer core is about 7-1 to about 20-1 and where the LHRH analog is p-Glu-His-Trp-Ser-Tyr-D-Trp-Leu-Arg-Pro-Gly-NH₂.

28. A bound microparticle according to claim 6 wherein the peptide is an LHRH analog.

29. A bound microparticle according to claim 28 wherein the ratio of glycolate units to tartrate residues

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is about 7-1 to about 20-1 and the LHRH analog is p-Glu-His-Trp-Ser-Tyr-D-Trp-Leu-Arg-Pro-Gly-NH₂.

30. A bound microparticle according to claim 4 wherein the peptide is a somatostatin analog.

5 31. A bound microparticle according to claim 30 wherein the ratio of glycolate units to citrate residues is about 7-1 to about 20-1 and the somatostatin analog is H-β-D-Nal-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH₂, where the two Cys residues are bonded by a disulfide bond, N-hydroxyethylpiperazinyl-acetyl-D-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂, where the two Cys residues are bonded by a disulfide bond or N-hydroxyethylpiperazinyl-ethylsulfonyl-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂, where the two Cys residues are bonded by a disulfide bond.

15 32. A bound microparticle according to claim 6 wherein the peptide is a somatostatin analog.

33. A bound microparticle according to claim 32 wherein the ratio of glycolate units to tartrate residues is about 7-1 to about 20-1 and the somatostatin analog is 20 H-β-D-Nal-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH₂, where the two Cys residues are bonded by a disulfide bond, N-hydroxyethylpiperazinyl-acetyl-D-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂, where the two Cys residues are bonded by a disulfide bond or N-hydroxyethylpiperazinyl-ethylsulfonyl-25 Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂, where the two Cys residues are bonded by a disulfide bond.

34. An encased microparticle comprising one or more bound microparticles according to claim 26 encased within an absorbable encasing polymer which comprises (a) 1-lactide based units and glycolide based units, 30 (b) d,l-lactide based units and glycolide based units, (c) d,l-lactide based units or (d) l-lactide based units and d,l-lactide based units.

35 35. An encased microparticle according to claim 34 wherein the ratio of glycolate units to citrate residues of the absorbable polymer core is about 7-1 to about 20-1, the

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LHRH analog is p-Glu-His-Trp-Ser-Tyr-D-Trp-Leu-Arg-Pro-Gly-NH₂ and where the ratio of:

- (a) l-lactide based units to glycolide based units is about 75-25 to about 90-10,
- 5 (b) d,l-lactide based units to glycolide based units is about 75-25 to about 90-10 and
- (c) l-lactide based units to d,l-lactide based units is about 80-20.

36. An encased microparticle comprising one or more
- 10 bound microparticles according to claim 28 encased within an absorbable encasing polymer which comprises (a) l-lactide based units and glycolide based units,
- (b) d,l-lactide based units and glycolide based units,
- (c) d,l-lactide based units or
- 15 (d) l-lactide based units and d,l-lactide based units.

37. An encased microparticle according to claim 36 wherein the ratio of glycolate units to tartrate residues of the absorbable polymer core is about 7-1 to about 20-1, the LHRH analog is p-Glu-His-Trp-Ser-Tyr-D-Trp-Leu-Arg-Pro-Gly-NH₂ and where the ratio of:
- 20 (a) l-lactide based units to glycolide based units is about 75-25 to about 90-10,

- (b) d,l-lactide based units to glycolide based units is about 75-25 to about 90-10 and
- 25 (c) l-lactide based units to d,l-lactide based units is about 80-20.

38. An encased microparticle comprising one or more bound microparticles according to claim 30 encased within an absorbable encasing polymer which comprises
- 30 (a) l-lactide based units and glycolide based units,
- (b) d,l-lactide based units and glycolide based units,
- (c) d,l-lactide based units or
- (d) l-lactide based units and d,l-lactide based units.

39. An encased microparticle according to claim 38 wherein the ratio of glycolate units to citrate residues of the absorbable polymer core is about 7-1 to about 20-1, the somatostatin analog is H- β -D-Nal-Cys-Tyr-D-Trp-Lys-Val-
- 35

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10 (b) d,l-lactide based units to glycolide based units is
about 75-25 to about 90-10 and
(c) l-lactide based units to d,l-lactide based units is
about 80-20.

41. An encased microparticle according to claim 40 wherein the ratio of glycolate units to tartrate residues of the absorbable polymer core is about 7-1 to about 20-1, the somatostatin analog is H- β -D-Nal-Cys-Tyr-D-Trp-Lys-Val-Cys-Thr-NH₂, where the two Cys residues are bonded by a disulfide bond, N-hydroxyethylpiperazinyl-acetyl-D-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂, where the two Cys residues are bonded by a disulfide bond or N-hydroxyethylpiperazinyl-ethylsulfonyl-Phe-Cys-Tyr-D-Trp-Lys-Abu-Cys-Thr-NH₂, where the two Cys residues are bonded by a disulfide bond; and where the ratio of:

(a) l-lactide based units to glycolide based units is about 75-25 to about 90-10,
(b) d,l-lactide based units to glycolide based units is
35 about 75-25 to about 90-10 and
(c) l-lactide based units to d,l-lactide based units is about 80-20.

43. A process according to claim 42 wherein a
5 dispersion of said bound microparticles in a solution
comprising said absorbable encasing polymer and a solvent
is dropped onto a pre-cooled medium, where said medium is
not a solvent of said absorbable encasing polymer.

44. A process according to claim 43 wherein the
10 solution of the absorbable encasing polymer consists of
about 5% to 30% of the absorbable encasing polymer, the
pre-cooled medium is an alcohol having two or more carbon
atoms and the temperature of the medium is room temperature
to about -80°C.

15 45. A process according to claim 44 wherein the temperature of the pre-cooled medium is about -60°C to -80°C and the medium is isopropyl alcohol.

46. A process for making an encased microparticle according to claim 12 comprising the step of encasing a bound microparticle with an absorbable encasing polymer using an emulsion technique.

[illegible]

COMBINED DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled ABSORBABLE MICROPARTICLES, the specification of which:

☐ is attached hereto.

☒ was filed on July 20, 2000 as Application Serial No. 09/600,648 and was amended on _____.

☒ was described and claimed in PCT International Application No. PCT/US99/01180 filed on 20 January 1999, and as amended under PCT Article 19 on _____.

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose all information I know to be material to patentability in accordance with Title 37, Code of Federal Regulations, §1.56.

I hereby claim the benefit under Title 35, United States Code, §119(e)(1) of any United States provisional application(s) listed below:

U.S. Serial No.	Filing Date	Status

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose all information I know to be material to patentability as defined in Title 37, Code of Federal Regulations, §1.56(a) which became available between the filing date of the prior application and the national or PCT international filing date of this application:

U.S. Serial No.	Filing Date	Status
09/015,394	29 January 1998	Abandoned

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I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign application(s) for patent or inventor's certificate or of any PCT international application(s) designating at least one country other than the United States of America listed below and have also identified below any foreign application for patent or inventor's certificate or any PCT international application(s) designating at least one country other than the United States of America filed by me on the same subject matter having a filing date before that of the application(s) of which priority is claimed:

Country	Application No.	Filing Date	Priority Claimed
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I hereby appoint the following attorneys and/or agents to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patents issued thereon.

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